## **AMENDMENTS**

This listing replaces all prior versions and listings of claims in the application.

- 1. (Currently amended) A stable liquid medical formulation, which contains (A) that comprises a therapeutically effective amount of an antibody in a glutamate buffer against CD40, sorbitol as isotonizing agent, a polysorbate as surfactant and glutamate as sole buffer and (B) that has a pH between 4.0 and 6.0.
- 2. (Currently amended) The <u>stable</u> liquid medical formulation according to claim 1, wherein the concentration of the buffer is between 1 mM and 50 mM.
  - 3. (Canceled)
- 4. (Currently amended) The <u>stable</u> liquid medical formulation according to claim [[3]] 1, which contains no salt as an isotonizing agent.
  - 5-6. (Canceled)
- 7. (Currently amended) The <u>stable</u> liquid medical formulation according to <del>any one</del> of claims 3 to 6 claim 1, wherein the <u>having an</u> osmotic pressure [[is]] between 250 mOsm and 350 mOsm.
  - 8. (Canceled)
- 9. (Currently amended) The <u>stable</u> liquid medical formulation according to claim [[8]] 1, wherein the surfactant is polysorbate 80.
- 10. (Currently amended) The <u>stable</u> liquid medical formulation according to claim § or 9 1, wherein the concentration of the surfactant [[is]] is present in a concentration between 0.02 mg/mL and 0.10 mg/mL.
- 11. (Currently amended) The <u>stable</u> liquid medical formulation according to claim 1, wherein the antibody <u>against CD40</u> is a human antibody, a humanized antibody, or a chimeric antibody.

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- 12. (Currently amended) The <u>stable</u> liquid medical formulation according to claim 1, wherein the antibody <u>against CD40</u> is a monoclonal antibody.
- 13. (Currently amended) The <u>stable</u> liquid medical formulation according to claim 1, wherein the antibody <u>against CD40</u> is IgG.
- 14. (Currently amended) The <u>stable</u> liquid medical formulation according to claim 13, wherein the IgG <u>subclass</u> is any one of IgG1, IgG2, or IgG4.
  - 15-17. (Canceled)
- 18. (Currently amended) The <u>stable</u> liquid medical formulation according to claim 1, wherein the antibody <u>against CD40</u> is present in a concentration between approximately 1 and 200 mg/mL.

19-20. (Canceled)

- 21. (Currently amended) The <u>stable</u> liquid medical formulation <del>according to any one</del> of claims 1, 19 or 20, which contains comprising:
  - (a) a therapeutically effective amount of an antibody against CD40;
  - (b) sorbitol as isotonizing agent;
  - (c) a polysorbate as surfactant; and
- (d) at least one stabilizing agent selected from the group consisting of glycine, methionine, cysteine hydrochloride, leucine, lysine hydrochloride, arginine hydrochloride, aspartic acid, ascorbic acid, EDTA, and salts thereof, with glutamate as sole buffer, the formulation having a pH between 4.0 and 6.0.

22-23. (Canceled)